- 2. (Amended) [Use according to claim] The method of Claim 1 or 20, wherein [the influence is an activation of] said modulating is an activation of blood vessel formation.
- 3. (Amended) [Use of a tissue factor or a fragment thereof for influencing the] A method for enhancing wound healing in a subject in need, comprising [by induction of a local expression of a tissue factor nucleic acid or by local application of] locally administering a functional tissue factor protein in a therapeutically effective amount to said subject in need.
- 4. (Amended) [Use according to claim] The method of Claim 3 or 21, wherein the said subject in need is afflicted with [wound healing in the case of] diabetis mellitus, vasculitis, arterial conclusive disease, chronic venous and infected ulcer, innervation impairment, decubital gangrene [and] or weak sutures [in the case of operations is concerned] after a surgery.
- 5. (Amended) [Use according to claim 1 or] The method of Claim [1 or] 2, wherein [the blood vessel formation in the case of] said subject in need is afflicted with arteriosclerosis, Crohn's disease [and], ulcerative colitis, diabetic retinopathy [and], or deep venous thrombosis of the legs/ulcus cruris [is concerned].
- 6. (Amended) [Use according to claim 1 or] The method of Claim 2, wherein the blood vessel formation [for replacing] is activated for the replacement of impaired blood vessels [is concerned].

- 2, wherein the tissue factor or a fragment thereof is present as expressible nucleic acid.
 - 8. (Amended) [Use according to claim] The method of Claim 7, wherein [the expression of the] said nucleic acid is [transient] expressed transiently.
 - 9. (Amended) [Use according to claim] The method of Claim 7 [or 8], wherein [the] said nucleic acid is a DNA.
 - 10. (Amended) [Use according to any one of claims] The method of Claim 7 [to 9], wherein [the] said nucleic acid is controlled by a constitutive or an indicuble promoter.
 - 11. (Amended) [Use according to any one of claims] The method of Claim 7 [to 10], wherein the nucleic acid is present in a Sindbis virus replicon vector.
 - 12. (Amended) [Use according to any one of claims] The method of Claim 7 [to 10], wherein the nucleic acid is controlled by a CMV or 5V40 promoter.
 - 13. (Amended) [Use according to any one of claims] The method of Claim 1 [to 12], 3, 20, or 21, wherein the tissue factor is present in a liposome or on a carrier, particularly gold particle.

- 14. (Amended) [Use according to any one of claims 1] The method of Claim 13, wherein the tissue factor is present in combination with further factors promoting the formation of blood vessels.
- 15. (Amended) [Use according to claim] <u>The method of Claim</u> 14, wherein [the] said further factors are present as expressible nucleic acids or functional proteins.
- 16. (Amended) [Use according to claim 14 or] <u>The method of Claim 15</u>, wherein [one of the factors] <u>at least one further factor is present and said further factor is VEGF.</u>
- 17. (Amended) [Use according to any one of claims] The method of Claim 1 [to 16], 3, 20, or 21, wherein the tissue factor is present in a pharmaceutical composition.

Please add the following new Claims 18-21:

- 18. (New) A pharmaceutical composition for modulation of blood vessel formation, comprising tissue factor or a fragment thereof and a pharmaceutically acceptable carrier.
- 19. (New) A pharmaceutical composition for activation of blood vessel formation, comprising tissue factor or a fragment thereof and a pharmaceutically acceptable carrier.